

# **Technical Information**

#### **Nitrofurantoin Broth Base**

**Product Code: DM 1857** 

Application: - Nitrofurantoin Broth Base is recommended d for the selective enrichment and isolation of Pseudomonas species.

### Composition\*\*

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Ingredients	Gms / Litre	
Peptic digest of animal tissue	7.500	
Casein enzymic hydrolysate	7.500	
Sodium chloride	5.000	
Final pH ( at 25°C)	7.2±0.2	

<sup>\*\*</sup>Formula adjusted, standardized to suit performance parameters

### Principle & Interpretation

Selective and differentiating media consisting of simple chemical components have been developed, both in solid and in liquid form, for culturing *Pseudomonas aeruginosa*. Nitrofurantoin, in the form of Macrodantin, has been shown to be active against most strains of *Escherichia coli, Staphylococcus aureus* and *Enterococcus faecalis* both in vitro and in clinical infections. Nitrofurantoin is not active against most strains of *Proteus* species or *Serratia* species. It has no activity against *Pseudomonas* species (1). Therefore nitrofurantoin incorporated in medium can be used as a selective medium for culturing of *Pseudomonas* species.

Casein enzymic hydrolysate and peptic digest of animal tissue supply the essential nutrients especially nitrogenous sources. Nitrofurantoin, 1-[(5-nitrofurfurylidene) amino] hydantoin, is a synthetic antibacterial agent which is effective against most common gram-negative and gram-positive urinary tract pathogenic bacteria (2).

### Methodology

Suspend 20 grams of dehydrated powder media in 1000 ml distilled water. Mix thoroughly & heat if necessary to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to room temperature and aseptically add 50 ml sterile 0.2% nitrofurantoin solution. Shake well and dispense in tubes or flasks as desired. Sterile nitrofurantoin solution (0.2%) is prepared by dissolving 1 gm Nitrofurantoin in 500 ml polyethylene glycol 300.

Note: Autosterilization takes place in 3 months. This solution can be stored for 6 months or longer.

## **Quality Control**

#### Appearance

Cream to yellow homogeneous free flowing powder

#### **Colour and Clarity**

With added nitrofurantoin: Fluorescent yellow coloured clear solution without any precipitate

#### Reaction

Reaction of 2.0% w/v aqueous solution at 25°C. pH: 7.2±0.2

#### pH Range

7.00-7.40

#### **Cultural Response**

DM 1857: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.





Organism	Inoculum (CFU)	Growth
Escherichia coli ATCC 25922	>=10³	inhibited
Pseudomonas aeruginosa ATCC 27853	50-100	good-luxuriant
Staphylococcus aureus ATCC 25923	>=103	inhibited

## Storage and Shelf Life

**Dried Media:** Store below 30°C in tightly closed container and the prepared medium at 2 - 8°C. Use before expiry date on the label. **Prepared Media:** 2-8°in sealable plastic bags for 2-5 days.

## **Further Reading**

- 1. Clinical and Laboratory Standards Institute, 2006, Performance standards for antimicrobial susceptibility testing. Approved standard M100-S15, Vol. 25, CLSI, Villanova, Pa.
- 2. Chamberlain R. E., 1976, Chemotherapeutic properties of prominent nitrofurans, J. Antimicrob. Chemother. 2:325336

### Disclaimer:

- User must ensure suitability of the product(s) in their application prior to use.
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